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(54) CANNABIS PLANT NAMED 'BIHEMP 050924'

(50) Latin Name: *Cannabis* hybrid Varietal Denomination: **BIHEMP 050924**

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 CPC ... A01H 5/12; A01H 5/02; A01H 5/00; A01H 6/28; A01H 6/00 See application file for complete search history.

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(57) ABSTRACT

The present invention provides a new and distinct *cannabis* cultivar designated as 'BIHEMP **050924**.' Disclosed herein are main terpenes of 'BIHEMP **050924**,' which are beta-caryophyllene, limonene, alpha-humulene, linalool, myrcene, trans-ocimene, beta-pinene, fenchol, and alpha-terpineol. Also, the present invention provides the estimated concentration of the THC $_{max}$, CBD $_{max}$, and CBG $_{max}$, about 0.21-0.43%, about 5.02-10.86%, and about 0.10-0.72%, respectively, at the time of assaying metabolites from flower samples of 'BIHEMP **050924**.'

7 Drawing Sheets

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Latin name of genus and species: *Cannabis* hybrid. Variety denomination: 'BIHEMP 050924'.

BACKGROUND OF THE INVENTION

The present invention relates to a new and distinct *can-nabis* cultivar designated as 'BIHEMP 050924'.

This new cultivar is the result of controlled-crosses between proprietary cultivars made by the inventors. The 10 new cultivar of 'BIHEMP 050924' was asexually reproduced via a stem 'cutting' and 'cloning' method by the inventors at Salinas, Calif. Asexual clones from the original source have been tested in greenhouses, nurseries, and/or fields. The properties of each cultivar were found to be 15 transmissible by such asexual reproduction. The cultivar is stable and reproduces true to type in successive generations of asexual reproduction.

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TAXONOMY AND NOMENCLATURE

Cannabis, more commonly known as marijuana, is a genus of flowering plants that includes at least three species, Cannabis sativa, Cannabis indica, and Cannabis ruderalis as determined by plant phenotypes and secondary metabolite profiles. In practice however, cannabis nomenclature is often used incorrectly or interchangeably. Cannabis literature can be found referring to all cannabis varieties as "sativas" or all cannabinoid producing plants as "indicas". Indeed the promiscuous crosses of indoor cannabis breeding programs have made it difficult to distinguish varieties, with most cannabis being sold in the United States having features of both sativa and indica species.

Human cultivation history of *Cannabis* dates back 8000 years (Schultes, R E., 1970, Random thoughts and queries on the botany of *Cannabis*. Pages 11-38 in: CRB Joyce, and S H Curry eds., THE BOTANY AND CHEMISTRY OF

CANNABIS. J. & A. Churchill. London, England). Hemp cloth recovered in Europe dates back 6000 years (Small, E, Beckstead, H D, and Chan, A, 1975, The evolution of cannabinoid phenotypes in Cannabis, ECONOMIC BOTANY 29(3):219-232). The written record of the pharmacologic properties of Cannabis goes back more than 4000 years (Ti, H. 2737 BC. NEI JING SU WEN HUANG TI, Yellow Emperor's Classic on Internal Medicine; referred to without citation in Small et al. 1975 Supra).

The taxonomy and nomenclature of the highly variable genus Cannabis (Emboden, W A, 1974, ECONOMIC BOTANY 28(3):304-310; Small, E and Cronquist, A, 1976, TAXON 25(4):405-435; Small E and Cronquist, A, 1977, TAXON 26(1):110; Hillig, K W and Mahlberg, P G, 2004, 15 American Journal of Botany 91(6):966-975), remains in question. This is in spite of the fact that its formal scientific name, 'Cannabis sativa L.', assigned by Carolus Linneaus (Linnaeus, C. 1753, SPECIES PLANTARUM, 2:1027, Salvius, Stockholm, Facsimile edition, 1957-1959, Ray Society, 20 London, U.K.), is one of the oldest established names in botanical history and is still accepted to this day. Another species in the genus, 'Cannabis indica Lam.' was formally named somewhat later (de Lamarck, JB, 1785, ENCYCLO-PEDIE METHODIQUE DE BOTANIQUE, 1(2):694-695), 25 characteristics of the plants. but is still very old in botanical history. In 1785, Jean-Baptiste Lamarck published a description of a second species of Cannabis, which he named Cannabis indica. Lamarck based his description of the newly named species on plant specimens collected in India. C. indica was described as relatively short, conical, and densely branched, whereas C. sativa was described as tall and laxly branched (Schultes R. E. et al, 1974, Harvard University Botanical Museum Leaflets, 23:337-367). C. indica plants were also described as having short, broad leaflets whereas those of C. 35 sativa were characterized as relatively long and narrow (Anderson L. C., 1980, Harvard University Botanical Museum Leaflets, 28:61-69). C. indica plants conforming to Schultes' and Anderson's descriptions may have originated from the Hindu Kush mountain range. Because of the often 40 harsh and variable (extremely cold winters, and warm summers) climate of those parts, C. indica is well-suited for cultivation in temperate climates.

Three other species names were proposed in the 1800s to distinguish plants with presumably different characteristics (C. macrosperma Stokes, C. chinensis Delile, C. gigantean Vilmorin), none of which are accepted today, although the epithet "indica" lives on as a subspecies of C. sativa ('C. sativa ssp. indica Lam.', Small and Cronquist 1976 Supra).

In the 20th century, two new names were added to the 50 liturgy of proposed 'Cannabis species: C. ruderalis' Janischevsky and a hybrid, x 'C. intersita' Sojak. (Small, E, Jui, P Y, and Lefkovitch, L P, 1976, SYSTEMATIC BOTANY 1(1):67-84; Small and Cronquist 1976 Supra). Further, numerous names have been proposed for horticultural vari- 55 ants of 'Cannabis' but as of 1976, "very few of these have been validly published as formal taxa under the International Code of Botanical Nomenclature" (Small and Cronquist 1976 Supra). Moreover, other recent work continues to focus on higher-order evolutionary relationships of the genus. 60 Cannabis has been variously ascribed as belonging to mulberry family (Moraceae) (Engler, H G A, Ulmaceae, Moraceae and Urticaceae, pages 59-118 in: A. Engler and K. Prantl eds., 1889, DIE NATURLICHEN PFLANZENFAMI-LIEN 3(1). W. Engelmann, Leipzig, Germany; Judd, W S, 65 Sanders, R W, and Donoghue, M J, 1994, HARVARD

PAPERS IN BOTANY 5:1-51; Humphries, C J and Blackmore, S, A review of the classification of the Moraceae, pages 267-277 In: Crane and Blackmore 1989 id.); nettle family (Urticaceae) (Berg, C C, Systematics and phylogeny of the Urticales, pages 193-220, in: P. R. Crane and S. Blackmore eds., 1989, EVOLUTION, SYSTEMATIC, AND FOSSIL HISTORY OF THE HAMAMELIDAE, VOL. 2, HIGHER HAMAMELIDAE, Clarendon Press, Oxford, U.K.); and most recently in its own family with hops (Humulus), Cannabaceae, or hemp family (Sytsma, K J, et al, 2002, AMERICAN JOURNAL OF BOTANY 89(9): 1531-1546). While the work of Small and Cronquist 1976 Supra, seemed to effectively confine the genus to a single species with 2 subspecies (C. sativa s., C. s. indica), each with two varieties (C. s. s. var. sativa, C. s. s. var. spontanea; C. s. i. var. indica, C. s. i. var. Kafiristanica) largely on the basis of chemotaxonomy and interfertility of all forms, more recent work (Sytsma et al. 2002 Supra), proposes a twospecies concept, resurrecting the binomial C. indica Lam. Since Sytsma et al. (2002) provides no key for discriminating between the species, the dichotomous key of Small and Cronquist (1976), which accounts for all forms in nature, whether wild or domesticated, is preferred to classify the

BRIEF SUMMARY OF THE INVENTION

This invention relates to a new and distinctive *cannabis* cultivar designated as 'BIHEMP 050924'.

The objective of the breeding program which produced novel plants disclosed herein was primarily to develop a *cannabis* cultivar with its unique blend of various cannabinoids and/or terpenes for (a) medicinal effects such as improving appetite and reducing nausea, vomiting and/or chronic pain, as well as neurological and cardiovascular effects, (b) psychoactive effects such as increased motivation and energetic behavior rather than indifference, passiveness and lethargy, and (c) recreational effects with enhanced enjoyment such as food and aroma.

As used herein, the term "cultivar" is used interchangeably with "variety", "strain", and/or "clone".

Cannabis plants produce a unique family of terpenophenolic compounds. Cannabinoids, terpenoids, and other compounds are secreted by glandular trichomes that occur most abundantly on the floral calyxes and bracts of female plants. As a drug it usually comes in the form of dried flower buds (marijuana), resin (hashish), or various extracts collectively known as hashish oil. The cannabis plant has at least 545 distinct compounds that span 20 chemical classes including cannabinoids, terpenes, terpenoids, amino acids, nitrogenous compounds, simple alcohols, aldehydes, ketones, esters, lactones, acids, fatty acids, steroids, noncannabinoid phenols, pigments, flavonoids, vitamins, proteins, enzymes, glycoproteins, and hydrocarbons. Terpenes and/or cannabinoids, in particular, have shown great potential in terms of medicinal value.

Terpenes and/or cannabinoids have been shown to be largely responsible for beneficial effects of a *cannabis* plant. In fact, each *cannabis* plant has the varying concentrations of medically viable compounds depending on different strains (genotypes) and their resulting chemotypes. Even a small variation in terpene and/or cannabinoid concentration can cause noticeable differences in the entourage and/or synergistic effects of a *cannabis* plant, which distinguishes

one variety from another. Research shows that it relies heavily on the physiological effects produced by terpenes and/or cannabinoids.

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Over 100 different kinds of terpenes have been identified in *cannabis* plants although not being as well-studied as ⁵ cannabinoids, they are instrumental in giving rise to the physiological and psychoactive effects in *cannabis*.

Terpenes are a large and diverse class of organic compounds, produced by a variety of plants. They are often strong smelling and thus may have had a protective function. Terpenes are an important component, not only influencing taste and smell of each cannabis strain but also influencing its effects on the mind and body of a subject such as humans and animals. Terpenes are a classification of organic molecules that are found in a wide variety of plants and animals. These molecules are known for their characteristic scents and flavors. The varying terpene concentrations found in cannabis plants directly influence the resulting taste and smell, as well as the observed effects. Non-limiting 20 examples of terpenes include Hemiterpenes, Monoterpenes, Sesquiterpenes, Diterpenes, Sesterterpenes, Triterpenes, Sesquarterpenes, Tetraterpenes, Polyterpenes, and Norisoprenoids. The main terpenes found in cannabis plants include, but are not limited to, myrcene, limonene, 25 caryophyllene, pinene, terpinene, terpinolene, camphene, terpineol, phellandrene, carene, humulene, pulegone, sabinene, geraniol, linalool, fenchol, borneol, eucalyptol, and nerolidol.

Cannabinoids are the most studied group of the main 30 physiologically active secondary metabolites in cannabis. The classical cannabinoids are concentrated in a viscous resin produced in structures known as glandular trichomes. At least 113 different cannabinoids have been isolated from 35 cannabis plants. The main classes of cannabinoids from cannabis include tetrahydrocannabinol (THC), cannabidiol (CBD), cannabigerol (CBG), and cannabinol (CBN). Cannabinoid can be at least one of a group comprising tetrahydrocannabinol (THC), cannabidiol (CBD), cannabigerol 40 (CBG), cannabinol (CBN) cannabichromene (CBC), cannabinodiol (CBDL), cannabicyclol (CBL), cannabivarin (CBV), tetrahydrocannabivarin (THCV), cannabidivarin (CBDV), cannabigerovarin (CBGV), cannabichromevarin (CBCV), cannabigerol monomethyl ether (CBGM), canna- 45 bielsoin (CBE), cannabicitran (CBT), cannabinol propyl variant (CBNV), cannabitriol (CBO), tetrahydrocannabinolic acid (THCA), tetrahydrocannabivarinic acid (THCVA), cannabidiolic acid (CBDA), cannabigerolic acid (CBGA) and cannabinerolic acid.

Most cannabinoids exist in two forms, as acids and in neutral (decarboxylated) forms. The acidic form of cannabinoids is designated by an "A" at the end of its acronym (i.e. THCA). The cannabinoids in their acidic forms (those ending in "-A") can be converted to their non-acidic forms through a process called decarboxylation when the sample is heated. The phytocannabinoids are synthesized in the plant as acidic forms. While some decarboxylation does occur in the plant, it increases significantly post-harvest and the kinetics increase at high temperatures (Flores-Sanchez and Verpoorte, 2008, Plant Cell Physiol. 49(12): 1767-1782). The biologically active forms for human consumption are the neutral forms. Decarboxylation is usually achieved by thorough drying of the plant material followed by heating it, often by combustion, vaporization, heating, or baking in an

oven. Unless otherwise noted, references to cannabinoids in a plant include both the acidic and decarboxylated versions (e.g., CBD and CBDA).

The molecules lose mass through the process of decarboxylation. In order to find the total theoretical active cannabinoids, the acid forms should be multiplied by 87.7%. For example, THCA can be converted to active THC using the formula: THCA×0.877=THC. The maximum THC for the sample is: THC $_{max}$ =(THCA×0.877)+THC. This method has been validated according to the principles of the International Conference on Harmonization. Similarly, CBDA can be converted to active CBD and the yield is determined using the yield formula: CBDA×0.877=CBD. Also the maximum amount of CBD yielded, i.e. max CBD for the sample is: CBD $_{max}$ =(CBDA×0.877)+CBD. Additionally, CBGA can be converted to active CBG by multiplying 87.8% to CBGA. Thus, the maximum amount of CBG is: CBG $_{max}$ =(CBGA×0.878)+CBG.

The biologically active chemicals found in plants, phytochemicals, may affect the normal structure or function of the human body and in some cases treat disease. The mechanisms for the medicinal and psychoactive properties of a *cannabis* plant, like any medicinal herb, produce the pharmacologic effects of its phytochemicals, and the key phytochemicals for a medical *cannabis* plant are cannabinoids and terpenes.

Δ9-Tetrahydrocannabinol (THC) is a psychoactive cannabinoid responsible for many of the effects such as mild to moderate pain relief, relaxation, insomnia and appetite stimulation. THC has been demonstrated to have anti-depressant effects. The majority of strains range from 12-21% THC with very potent and carefully prepared strains reaching even higher. While Δ9-Tetrahydrocannabinol (THC) is also implicated in the treatment of disease, the psychotropic activity of THC makes it undesirable for some patients and/or indications.

Tetrahydrocannabinol, THC, is the primary psychoactive and medicinal cannabinoid and is the result of the decarboxylation of tetrahydrocannabinolic acid (THC-A), its acidic precursor. THC-A, (6ar,10ar)-1-hydroxy-6,6,9-trimethyl-3-pentyl-6a,7,8,10a-tetrahydro-6h-benzochromene-2-carboxylic acid, is found in the trichomes of the plant and converted into THC, which actually exists in only minute quantities in the living plant, after harvest and drying.

Cannabidiol (CBD) is one of the principal cannabinoids found in a cannabis plant and is largely considered to the most medically significant. CBD occurs in many strains, at low levels, <1%. In some cases, CBD can be the dominant cannabinoid, as high as 15% by weight. CBD is nonpsychoactive, meaning that unlike THC, CBD does not cause a noticeable "high". CBD has shown potential for medical properties in the treatment of a wide variety of diseases and symptoms, including cancer, nausea, chronic pain, spasms, seizures/epilepsy, anxiety, psoriasis, Crohn's disease, rheumatoid arthritis, diabetes, schizophrenia, posttraumatic stress disorder (PTSD), alcoholism, strokes, multiple sclerosis, and cardiovascular disease. CBD also has been reported to act as a muscle relaxant, antibiotic, antiinflammatory, and bone stimulant, as well as to improve blood circulation, cause drowsiness, and protect the nervous system. It can provide relief for chronic pain due to muscle spasticity, convulsions and inflammation, as well as effective relief from anxiety-related disorders. It can offer relief for patients with Multiple Sclerosis (MS), Fibromyalgia and

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Epilepsy. CBD has also been shown to inhibit cancer cell growth when injected into breast and brain tumors in com-

A *cannabis* cultivar can be used to achieve the desire of patients to be treated with CBD without the adverse side- ⁵ effects (e.g., psychoactivity) of THC.

bination with THC.

Cannabichromene (CBC) is a rare, non-psychoactive cannabinoid, usually found at low levels (<1%) when present. It has been shown to have anti-depressant effects and to improve the pain-relieving effects of THC. Studies have demonstrated that CBC has sedative effects such as promoting relaxation.

Cannabidiol (CBD) and cannabichromene (CBC) are both non-psychoactive and end products of CBG metabolism, like THC, so that they can be used medically.

Cannabigerol (CBG) is a non-psychoactive cannabinoid. CBG-acid is the precursor to both THC-acid and CBD-acid in the plant usually found at low levels (<1%) when present. It has been demonstrated to have both pain relieving and inflammation reducing effects. CBG reduces intraocular pressure, associated with glaucoma. CBG has been shown to have antibiotic properties and to inhibit platelet aggregation, which slows the rate of blood clotting. While Cannabigerol (CBG), is not considered psychoactive, it is known to block the psychoactive effects of THC and is considered medically active in a variety of conditions. Its precursor, cannabigerolic acid, CBGA, (E)-3-(3,7-Dimethyl-2,6-octadienyl)-2,4-dihydroxy-6-pentylbenzoic acid, is being studied medically

Cannabinol (CBN) is an oxidative degradation product of THC. It may result from improper storage or curing and extensive processing, such as when making concentrates. It is usually formed when THC is exposed to UV light and oxygen over time. CBN has some psychoactive properties, 35 less strength than THC. CBN is thought to enhance the dizziness and disorientation that users of *cannabis* may experience. It may cause feelings of grogginess, but has been shown to reduce heart rate.

High potency *cannabis* plants contain large quantities of specific terpenes as well as various assortments of other terpenes. For instance, a *cannabis* plant may have a profile with either a high level of, a moderate amount of or a small amount of various terpenes depending on its cultivar and environmental conditions.

Various cultivars of 'Camabis' species have been cultivated in an effort to create a cultivar best suited to meet the interest of inventors according to their own need. The particular plant disclosed herein was discovered in the area where the inventors were intentionally cross-pollinating and cultivating plants described below using standard Mendelian breeding procedures well known to those of ordinary skill in the art. This resulted in the progenies of the inventors' crosses

The progenies resulting from any selection stage of either the crossing, selfing or backcrossing versions of the breeding regimes of the present invention were asexually reproduced to fix and maintain the desirable THC content, CBs content, terpenes content, the aroma and flavor(s) typical of the desired class, and the other desirable phenotypic and/or genotypic characteristics. The resultant selected *cannabis* cultivar is designated as 'BIHEMP 050924' disclosed herein.

The inventors reproduced progenies asexually by stem cutting and cloning. This is the origin of this remarkable new cultivar. The plant has been and continues to be asexually 8

reproduced by stem cutting and cloning at the inventors' greenhouses, nurseries and/or fields in Salinas, Calif., Oakland, Calif., and/or Washington, D.C.

The following are the most outstanding and distinguishing chemical characteristics of this new cultivar when grown under normal conditions in Salinas, Calif. Chemical analyses of the new *cannabis* variety and the check variety (or the parental varieties) disclosed herein were performed using standard chemical separation techniques well known to those skilled in the art. Samples for assaying were obtained from flower tissues of the *cannabis* plant disclosed herein. Cannabinoid composition of this cultivar can be determined by assaying the concentration of at least one cannabinoid in a subset (e.g., sample) of the harvested product.

Table 1 includes detailed information of the *cannabis* plant named 'BIHEMP 050924' including the concentration ranges of terpenes and cannabinoids as tested on flowers at least seven different times. The *cannabis* plant has been tested in a laboratory setting and/or facility to determine cannabinoids and terpenes concentrations in the *cannabis* plant named 'BIHEMP 050924' according to the procedures provided in Giese et al. (Journal of AOAC International (2015) 98(6):1503-1522).

- The main terpenes found in 'BIHEMP 050924' are beta-caryophyllene, limonene, alpha-humulene, linalool, myrcene, trans-ocimene, beta-pinene, fenchol, and alpha-terpineol; and
- 2) The estimated concentration of the total THC_{max} , CBD_{max} , and CBG_{max} is about 0.21-0.43%, about 5.02-10.86%, and about 0.10-0.72%, respectively, at the time of assaying metabolites from flower samples of 'BIHEMP 050924'.

Terpene and cannabinoid profiles of 'BIHEMP 050924' demonstrate that 'BIHEMP 050924' has a phenotypically unique profile, particular insofar as to the level of terpenes and cannabinoids. This data is presented in a tabular form in Table 1.

TABLE 1

F	Ranges of	Active Cann	abinoids a	and Terpenes	
R	anges of.	Active Canna	binoids (% by weight)	
Max THC	0.21- 0.43% Rang	Max CBD	10.86%	Max CBG weight)	0.10- 0.72%
thujene alpha-pinene	0.00%	gamma- terpinene linalool	0.00%	hexyl hexanoate octyl butyrate	0.00- 0.03% 0.00%
camphene	0.04% 0.00- 0.01%	oxide terpinolene	0.00- 0.02%	beta-caryo- phyllene	0.12- 0.60%
sabinene	0.00%	fenchone	0.00%	alpha- humulene	0.03- 0.38%
beta-pinene	0.01- 0.07%	linalool	0.02- 0.16%	cis-nerolidol	0.00%
myrcene	0.00- 0.12%	fenchol	0.01- 0.06%	trans-nerolidol	0.03- 0.05%
alpha- phellandrene	0.00%	_	_	caryophyllene oxide	0.00- 0.03%
carene	0.00%	camphor	0.00%	alpha-bisabolol	0.01- 0.03%
alpha-terpinene limonene	0.00% 0.06- 0.62%	isoborneol (-) borneol	0.00% 0.00- 0.02%	nerol geraniol	0.00% 0.00%
beta- phellandrene	0.00%	menthol	0.00%	geranyl-acetate	0.00%
cineole	0.00%	hexyl butyrate	0.00%	methyl- eugenol	0.00%

TABLE 1-continued

	Ranges of	`Active Canr	nabinoids	and Terpenes	
cis-ocimene	0.00%		0.01- 0.06%	Total Terpenes	0.61- 2.42%
trans-ocimene	0.00- 0.12%	citronellol	0.00%	_	_

The *cannabis* plant named 'BIHEMP 050924' has a complement of terpenes, including but not limited to, relatively high levels of beta-caryophyllene, limonene, alphahumulene, linalool, myrcene, trans-ocimene, beta-pinene, fenchol, and alpha-terpineol compared to other terpene compounds. This unique combination of differently concentrated terpenes further distinguishes 'BIHEMP 050924' from other varieties in its odor, its medical qualities, and its effects on mood and mentation.

Asexual Reproduction

As exual reproduction, also known as "cloning", is a $_{20}$ process well known to those of ordinary skill in the art of cannabis production and breeding and includes the following steps.

The *cannabis* cultivar disclosed herein is asexually propagated via taking cuttings of shoots and putting them in rock wool cubes. These cubes are presoaked with pH adjusted water and kept warm (~80° F.). Full trays are covered, left under 18 hours of light and allowed to root (7-14 days). Upon root onset, the plantlets are transplanted into rigid 1 gallon containers filled with a proprietary soil mix A and remain in 18 hours of daylight for another 14-21 days. Once root-bound, plants are transplanted into rigid 3 gallon containers filled with proprietary soil mix B. Immediately, the light cycle is altered to 12/12 and flower initiating begins. The plants remain in 12/12 lighting until harvesting. They undergo a propriety nutrient regimen and grow as undisturbed as possible for 60-70 days depending on chemotype analysis.

All sun leaves are removed and the plant is dismantled to result in approximately 12" branches covered in inflorescences and trichomes. The goal in harvesting is to actually harvest trichome heads but not 'buds'. Thus, great care is taken not to disturb the trichome heads and as much of the plant remains intact as possible to promote even and slow drying. Slow drying is followed by a one to two months 45 curing process.

Observation of the all female progenies of the original plant has demonstrated that this new and distinct cultivar has fulfilled the objectives and that its distinctive characteristics are firmly fixed and hold true from generation to generation vegetatively propagated from the original plant.

Under careful observation, the unique characteristics of the new cultivar have been uniform, stable and reproduced true to type in successive generations of asexual reproduction.

DESCRIPTION OF THE DRAWINGS

The accompanying color photographs depict characteristics of the new 'BIHEMP 050924' plants as nearly true as possible to make color reproductions. The overall appearance of the 'BIHEMP 050924' plants in photographs is shown in colors that may differ slightly from the color values described in the detailed botanical description.

FIG. 1 shows an overall view of the 'B̄IHEMP 050924' $_{65}$ plant from the side.

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FIG. 2A shows a close view of a single leaf of the check variety BLK03 plant.

FIG. 2B shows a close view of a single leaf of the new variety 'BIHEMP 050924' plant.

FIG. 3A shows top parts (including inflorescence) of the BLK03 plant from the side.

FIG. 3B shows top parts (including inflorescence) of the 'BIHEMP 050924' plant from the side.

FIG. 4 shows a close view of flowers of the 'BIHEMP 050924' plant at the late flowering/mature stage.

FIG. 5 shows another close view of flowers of the 'BIHEMP 050924' plant at the late flowering/mature stage.

DETAILED BOTANICAL DESCRIPTION

'BIHEMP 050924' has not been observed under all possible environmental conditions, and the phenotype may vary significantly with variations in environment. The following observations, measurements, and comparisons describe this plant as grown at Salinas, Calif., when grown in the greenhouse, nursery or field, unless otherwise noted.

Plants for the botanical measurements in the present application are annual plants. In the following description, the color determination is in accordance with The Royal Horticultural Society Colour Chart, 2007 Edition, except where general color terms of ordinary dictionary significance are used.

The *cannabis* plant disclosed herein was derived from female and male parents that are internally designated as below.

A GNBR internal Code of the cannabis plant named 'BIHEMP 050924' is 05.09.24. A GNBR Breeding Code of the cannabis plant named 'BIHEMP 050924' is (V24.S1.O3xV24.S1.N5.05)x(V24.S1.O3xV24.S1.N5.09) .24. The additional number '.24' was only assigned to the 24th individual plant (i.e. 'BIHEMP 050924') selected from progenies of the cross event between pollen acceptor (O3.N5.05) and pollen donor (O3.N5.09). 'BIHEMP 050924' is a fertile hybrid derived from a controlled-cross between two proprietary cultivars: V24.S1.O3xV24.S1.N5.05 (pollen acceptor; female parent), known as O3.N5.05 or 05 and V24.S1.O3xV24.S1.N5.09 (pollen donor; male parent), also known as O3.N5.09 or 09. The initial cross between two parental cultivars was made in April 2015. The primary phenotypic criteria used to select the new and distinct cannabis cultivar disclosed herein is as follows: structure score, nose/organoleptic testing, mold susceptibility/resistance, and insect susceptibility/resistance. Also, the first 55 asexual propagation of 'BIHEMP 050924' occurred on Mar. 25, 2017 in Salinas, Calif.

The following traits in combination further distinguish the *cannabis* cultivar 'BIHEMP 050924' from the check variety 'BLK03', which is set as a standard for phenotypic comparison. Tables 2 to 6 present phenotypic traits and/or characteristics of 'BIHEMP 050924' compared to the check variety 'BLK03' as follows. All plants were raised together and evaluated when 100 days old (i.e., 25 days in vegetative stage, 15 days in propagation stage, and 60 days in flowering times).

	General Characteris	rtice
Characteristics		
	New Variety	Check Variety (BLK03)
Plant life forms	An herbaceous plant (herb)	An herbaceous plant (herb)
Plant growth	An upright, tap-rooted	An upright, tap-rooted
habit	annual plant; forming	annual plant; forming
	fibrous roots when	fibrous roots
	asexually propagated	when asexually propagated
Plant origin	A controlled-cross	A controlled-cross between
	between pollen acceptor	pollen acceptor (GLD13)
	(O3.N5.05) and pollen donor (O3.N5.09)	and pollen donor (BSIA)
Plant	Asexually propagated	Asexually propagated
propagation	by stem	by stem
	cuttings and cloning	cuttings and cloning
Propagation	Easy	Moderate
ease Height	0.3-1.2 m	0.5-2.5 m
Width	56 cm	119.5 cm
Plant vigor	High	Medium
Time to	10 weeks	8 weeks
Harvest		
(Seed to Harvest)		
Resistance to	Resistant to pests as	Not Resistant to pests as
pests or	follows; (1)	follows; (1) two spotted
diseases	Western Flower Thrip	spider mite
	(Frankliniella	(Tetranychus urticae
	occidentalis); (2)	(Koch)); (2) Aphids
	Leaf Miner (<i>Liriomyza</i> sativae); (3) Whitefly	species such as: Cannabis Aphids (Phorodon
	(Trialeurodes vapor-	cannabis), Green Peach
	ariorum); (4) Lepidoptera	
	species such as:	(Sulzer)), Foxglove Aphid
	Armyworm (Spodoptera	(Aulacorthum
	frupperda), Cabbage Whites (Pieris rapae),	solani), Peach Aphid (Macrosiphum
	Painted Lady (Vanessa	euphorbiae) and Black
	cardui), Lepidoptera sp.	Bean whitefly
	Aphid (Aphis fabae); (3)	(Trialeurodes
	Resistant to diseases as	vaporariorum);
	follows; Botrytis/	(4) Lepidoptera species
	Flower Rot (Botrytis cinerea), Powdery	such as: Armyworm (Spodoptera frupperda),
	Mildew (Podosphaera	Cabbage Whites
	xanthii)	(Pieris rapae),
	•	Painted Lady (Vanessa
		cardui), Lepidoptera sp.
		Not resistant to diseases as follows; Botrytis/Flower
		Rot (Botrytis cinerea)
		and Powdery Mildew
		(Podosphaera xanthii)
Genetically-	NO	NO
modified		
organism		

TABLE 3

Leaf/Foliage		
Characteristics	New Variety	Check Variety (BLK03)
Leaf arrangement	Alternate	Alternate
Leaf shape	Palmately compound	Palmately compound
Leaf structure	Linear-lanceolate leaflet blades with glandular hairs	Linear-lanceolate leaflet blades with glandular hairs

TABLE 3-continued

		Leaf/Foliage	
	Characteristics	New Variety	Check Variety (BLK03)
	Leaf margins	Dentate, coarsely serrated, and the teeth point away	Dentate, coarsely serrated, and the teeth point away
)	Leaf hairs	from the tip Present on both upper and lower surfaces	from the tip Present on both upper and lower surfaces
	Leaf length with petiole at maturity	20.4 cm	16.6 cm
	Leaf width at maturity	9.8-14.6 cm	10.7 cm
5	Petiole length at maturity	7.7 cm	6.5 cm
	Petiole color (RHS No.)	53A at day 60 in flowering; 134B at day 5 in flowering	140C
	Intensity of petiole	Weak (early flowering	Medium (vegetativ
	anthocyanin	stage);	stage); very
)		Medium (late flowering stage - during days 30-60 in flowering)	strong (late flowering stage)
	Stipule length at maturity	0.6 cm	0.7 cm
	Stipule shape	Elliptical	Elliptical
	Stipule color (RHS No.)	134A	149B
5	No. of leaflets	5-7	5-7
,	Middle largest (longest) leaflet length	12.1 cm	9.8 cm
	Middle largest (longest) leaflet width	2.2 cm	2.3 cm
	Middle largest (longest) leaflet length/width ratio	12.1:2.2	9.8:2.3
)	No. teeth of middle leaflet (average)	22	25
	Leaf (upper side) color (RHS No.)	139B	132A
	Leaf (lower side) color (RHS No.)	139C	134D
5	Leaf glossiness	Strong at the upper surface	Strong
	Vein/midrib shape	Obliquely continuous throughout leaflet	Obliquely continuous throughout leaflet
	Vein/midrib color (RHS No.)	151D	144C
)	Aroma	Pungent, yet sweet	Spicy

TABLE 4

45		Stem	
	Characteristics	New Variety	Check Variety (BLK03)
50	Stem shape	Hollow, ribbed, large	Hollow, ribbed, textured
	Stem diameter at base	2.3 cm	2.8 cm
	Stem color (RHSNo.) Depth of main stem ribs/grooves	149D Shallow	N144D Absent
	Internode length	14.4-27.2 cm	2.4-4.9 cm

TABLE 5

	madel c	
Inflo	escence (Female/Pistillate	Flowers)
Characteristics	New Variety	Check Variety (BLK03)
Flowering (blooming) habit Proportion of female plants	Elongated compound cymes, from 0.2 m- 1.5 m in length 100% pistillate	Elongated compound cymes, from 0.5 m- 1.2 m in length 100% pistillate
	Characteristics Flowering (blooming) habit Proportion of female	Inflorescence (Female/Pistillate Characteristics New Variety Flowering Elongated compound (blooming) cymes, from 0.2 m-habit 1.5 m in length Proportion of female 100% pistillate

TABLE 5-continued

Inflo	Inflorescence (Female/Pistillate Flowers)			
Characteristics	New Variety	Check Variety (BLK03)		
Inflorescence position	Above	Even		
Flower arrangement	Cymose	Cymose (terminal bud matures, while lateral flowers mature thereafter)		
Number of flowers per plant	110-143 per cyme	80-120 per cyme		
Flower shape Flower (individual	Calcarate-urceolate 0.3 cm	Calcarate-urceolate 0.7 cm		
pistillate) length Flower (compound	4.7 cm	3 .8 cm		
cyme) diameter Corolla shape Corolla size	No defined corolla	No defined corolla		
Corolla Color (RHS No.)	n/a	n/a		
Bract shape Bract size	Urceolate 0.4-1.2 cm	Urceolate 0.2-0.8 cm		
Bract color (RHS No.)	130B	N134C		
Calyx shape Calyx color	No defined calyx n/a	No defined calyx n/a		
(RHS No.) Stigma shape	Linear	Acute		
Stigma length Stigma color	1.9 mm 53A	2.2 mm 159D		
(RHS No.) Trichome shape	Capitate-stalked glandula	r Capitate-stalked glandular		
Trichome color (RHS No.)	157A before harvest, at approximately day 38 of flowering	157A at day 40 in flowering		
Other types of trichomes	Capitate sessile tri- chomes are present on the leaves of plants, as well as being noticed in the flowers (color: 157A at day 38 in flowering). During later flowering, i.e. day 40 to day 60 in flowering, the capitate stalked	Capitate sessile tri- chomes are present on the leaves of plants, as well as being noticed in the flowers (color: 157A at day 40 in flowering). During later flowering, i.e. day 48 to day 60 in flowering, capitate stalked trichomes are		
	trichomes are present (color: N30B).	present (color: N30B). Bulbous and non- glandular trichomes are also present and most noticeable on the petioles, stems, and leaves (color: 157A).		
Terminal bud shape Terminal bud color (RHS No.)	Oblong 132C	Oblong 203C		
Pedicel	Absent	Absent		
Staminate shape	No staminate flowers produced naturally; however, male flower	No staminate flowers produced naturally; however, male flower		

(staminate) can be

induced with chemical

and silver thiosulphate

anionic complex).

compounds (such as

silver nitrate

(staminate) can be

induced with chemical

silver nitrate and silver

compounds (such as

thiosulphate anionic

complex).

TABLE 5-continued

14

5 Cl	naracteristics	New Variety	Check Variety (BLK03
	llen description	Absent	Absent
Se	ed shape	Smooth and globular	Smooth and globular
Se	ed size	2.0-2.8 mm	1.8-2.3 mm
M	arbling of seed	Weak	Absent (non-existent)
Pe	tal description	Apetalous	Apetalous
0 M	ax THC content	About 0.21-0.43%	About 18.88-19.37%
	ax CBD ntent	About 5.02-10.86 %	0.00%
	ax CBG ntent	About 0.10-0.72%	About 0.84-0.91 %

15 n/a: not available

TABLE 6

		Other Characteristics		
20	Characteristics	New Variety	Check Variety (BLK03)	
	Time period and condition of flowering/blooming	6-8 weeks	7-9 weeks	
25	Hardiness of plant Breaking action	Hardy to 25° Fambient temperature Flexible, highly resistant to breakage	Hardy to 25° Fambient temperature Strong, non-flexible	
30	Rooting rate after cutting/cloning	99%-vigorous	70%-moderate	
	Types of Cutting for Cloning	Stem	Stem	
	Shipping quality	Good	Moderate	
35	Storage life	Medium (2-5 months with minor changes in physical appearance and/or smell/taste)	Medium (2-6 months with minor changes in physical appearance and/ or smell/taste)	
	Market use Productivity of flower	Medicinal Approximately 0.136-0.363 kg	n/a Approximately 0.14-0.45 kg can	
40		can be produced per plant, dependent on finished plant size (1.0-2.6 m); Growing conditions/en- vironment will	be produced per plant; dependent on finished size (0.6-1.2 m); Growing conditions/en- vironment will	
45		dictate final yield/output	dictate final yield/output	

In general, 'BIHEMP 050924' is larger in height than both parents, (O3.N5.05) and (O3.N5.09). 'BIHEMP 050924' is more robust in terms of growing performance, time to rooted clones, and time to flower maturity. Also, 'BIHEMP 050924' has greater resistance to pests and diseases, stronger branches, thicker stems, greater flexibility, and higher yielding. 'BIHEMP 050924' clearly demonstrates hybrid vigor, and outperforms both parents overall. Chemically, 'BIHEMP 050924' has a higher propyl cannabinoid content as well as a higher terpene content than either parent.

When 'BIHEMP 050924' is compared to the check variety 'BLK03', 'BIHEMP 050924' is shorter in plant height and narrower in plant width than 'BLK03'. 'BIHEMP 050924' shows higher plant vigor and longer time to harvest than 'BLK03'. However, 'BIHEMP 050924' has longer leafs than 'BLK03' in terms of whole leaf length including petiole. Also, 'BIHEMP 050924' has longer leaflets than 'BLK03' when comparing the middle largest leaflet length, while 'BIHEMP 050924' has less teeth numbers in middle

leaflet than 'BLK03'. 'BIHEMP 050924' has a longer petiole but a little shorter stipule in average than 'BLK03' at maturity. Regarding the average stem diameter at base, 'BIHEMP 050924' is shorter than 'BLK03'. However, the internode length of 'BIHEMP 050924' is conspicuously 5 longer than that of 'BLK03'. In terms of flower numbers per cyme, 'BIHEMP 050924' has more flowers than 'BLK03'. When comparing the compound cyme diameter, 'BIHEMP 050924' is longer than 'BLK03', while individual pistillate flower of 'BIHEMP 050924' is shorter than that of 'BLK03'. 10 'BIHEMP 050924' has a longer bract than 'BLK03', while having a little shorter stigma. With respect to aroma, 'BIHEMP 050924' have a pungent yet sweet scent, while 'BLK03' has a generally spicy smell.

When 'BIHEMP 050924' is compared to the known 15 cannabis plant named 'ECUADORIAN SATIVA' (U.S. Plant Pat. No. 27,475), there are several distinctive characteristics. For example, overall form of 'BIHEMP 050924' plant is short in plant height but wider across at the widest point than the 'ECUADORIAN SATIVA' plant. 'BIHEMP 20 050924' plant has a little longer middle leaflet (without petiole) and whole leaf (with petiole) length than the 'ECUADORIAN SATIVA' plant. Generally, 'BIHEMP

050924' plant has a little shorter petiole at maturity than the 'ECUADORIAN SATIVA' plant. 'BIHEMP 050924' plant has a narrower middle leaflet width than the 'ECUADORIAN SATIVA' plant. Regarding stem diameter at base, 'BIHEMP 050924' is similar to 'ECUADORIAN SATIVA'. While the aroma of 'ECUADORIAN SATIVA' is strongly mephitic with hints of limonene, 'BIHEMP 050924' has a pungent yet sweet scent. When comparing total THC content between 'BIHEMP 050924' and 'ECUADORIAN SATIVA', the total THC content of 'BIHEMP 050924' is between 0.21-0.43%, while 'ECUADORIAN SATIVA' accumulates 12.45% total THC.

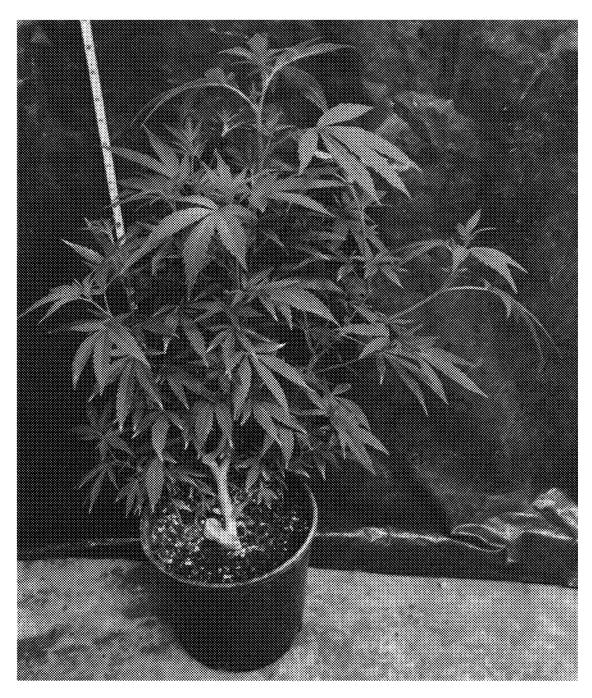
'BIHEMP 050924' having no more than 0.3% THC content, qualifies as a hemp under most growing conditions. As with other plants, 'BIHEMP 050924' may accumulate higher contents of total THC under some growing conditions, or if allowed to continue growing past maturity.

The invention claimed is:

1. A new and distinct cultivar of *Cannabis* plant named 'BIHEMP 050924' substantially as shown and described herein

* * * * *

Fig. 1



BIHEMP 050924

Fig. 2A



BLK03

Fig. 2B



BIHEMP 050924

Fig. 3A



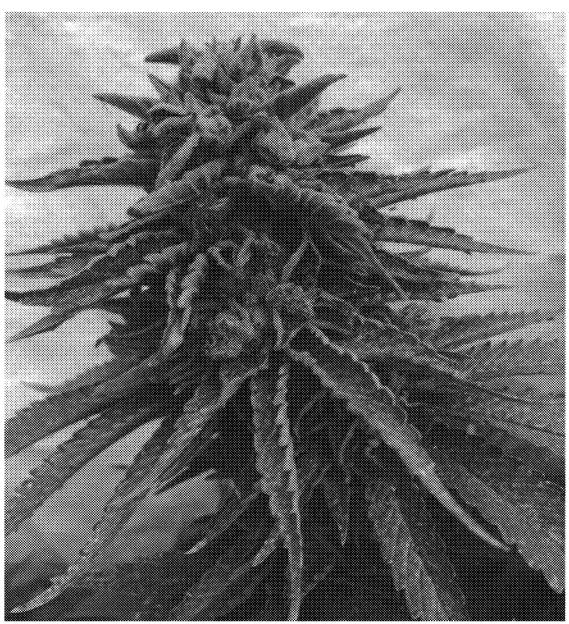
BLK03

Fig. 3B



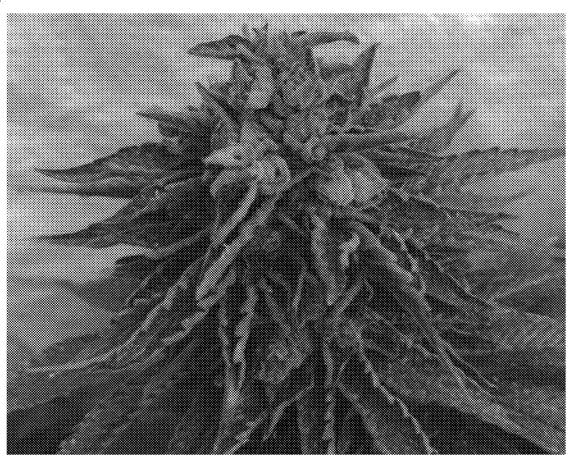
BIHEMP 050924

Fig. 4



BIHEMP 050924

Fig. 5



BIHEMP 050924